

This product is for research use only (not for diagnostic or therapeutic use)

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# Product no AS08 358

## Anti-SNCA | Alpha-synuclein

### **Product information**

Immunogen recombinant, full length, human alpha-synuclein UniProt: P37840, epitope was mapped between amino acid 1-15

**Host** Rabbit

Clonality Polyclonal

Purity Serum

Format Lyophilized

Quantity 100 μl

**Reconstitution** For reconstitution add 100 μl of sterile water

Storage Store lyophilized/reconstituted at -20°C; once reconstituted make aliquots to avoid repeated freeze-thaw cycles. Please remember to spin the tubes briefly prior to opening them to avoid any losses that might occur from material adhering to

the cap or sides of the tube.

**Additional information** Immunolocalization: human tissue was paraffin-embedded and sectioned. De-waxed and rehydrated in an ethanol gradient. Antigens were retrieved in sodium citrate buffer (pH 6) at 95 °C for 1 h. The tissue sections were separately incubated for 1 h at RT with primary antibody and antibody binding was visualized with IgG Preoxidase Reagent Kit.

> This antibody will recognize human SNCA monomers and multimers in Western blot and can be used to detect fibrills in a sandwich ELISA.

In ELISA this antibody can be used for detection, combined with AS13 2719 as a capture antibody.

# **Application information**

Recommended dilution 1:3000 (ELISA), 1-2 μg/ml (IL), 1:1000 (WB)

Expected | apparent 14 kDa

Confirmed reactivity Human

Predicted reactivity Primates, Chimpanzee, Gorilla

Selected references Bargar et al. (2021) Discrimination of MSA-P and MSA-C by RT-QuIC analysis of olfactory mucosa: the first assessment of assay reproducibility between two specialized laboratories. Mol Neurodegener. 2021 Dec 11;16(1):82.

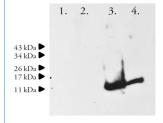
doi: 10.1186/s13024-021-00491-y. PMID: 34895275; PMCID: PMC8665327.

Brännström et al. (2014). A generic method for design of oligomer-specific antibodies. PLoS One. 2014 Mar

11;9(3):e90857. doi: 10.1371/journal.pone.0090857. eCollection 2014.

#### Western analysis

- Membrane: Nitrocellulose
  Blocking buffer: 5% dry milk in PBS, 0.15% tween 20
  Antibody dilution: 1:1000
  Secondary antibody: anti-rabbit (HRP)
  Detection: Enhanced Chemoluminescence (pico) 10s



- 1. Cell lysate lacking alpha-synuclein 10 000 ng (negative control)
- 2. Human plasma 10 000 ng (negative control)
- 3. Human alpha-synuclein 100 ng



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4. Human alpha-synuclien 10 ng